

Substitute for form 1449A/PTO

## **INFORMATION DISCLOSURE**

**STATEMENT BY APPLICANT**

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(use as many sheets as necessary)

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**COMPLETE IF KNOWN**

<b>Application Number</b>	09/937,499
<b>Filing Date</b>	February 7, 2002
<b>First Named Inventor</b>	Konstantin Petrukhin
<b>Group Art Unit</b>	1646
<b>Examiner Name</b>	
<b>Attorney Docket Number</b>	20430P

## **U.S. PATENT DOCUMENTS**

## **FOREIGN PATENT DOCUMENTS**

Examiner Signature	/Amy Juedes/	Date Considered	08/09/2006
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				Group Art Unit	1646
				Examiner Name	
Sheet	2	of	4	Attorney Docket Number	20430P

NON PATENT LITERATURE DOCUMENTS					
Examiner Initials*	Cite No.	Include name of the author, title, date, page(s), volume-issue number(s) and place of publication.			
AJ	B	Brian Robertson, "The real life of voltage-gated K+ channels: more than model behaviour", Trends in Pharmacol. Sci., 18:474-483 (December 1997)			
	C	Jan & Jan, "Voltage-gated and inwardly rectifying potassium channels", J. Physiology, 505.2:267-282 (1997)			
	D	Catterall, A. W., "Structure & Function of Voltage-Gated Ion Channels, Annual Rev. Biochem., 64:493-531 (1995)			
	E	Sullivan & Daiger, "Inherited retinal degeneration: exceptional genetic and clinical heterogeneity", Mol. Med. Today, 2:380-386 (September 1996)			
	F	Evans et al., "Genetic linkage of cone-rod retinal dystrophy to chromosome 19q and evidence for segregation distortion", Nature Genetics, 6:210-213 (February 1994)			
	G	Kelsell et al., "Localisation of a gene for dominant cone-rod dystrophy (CORD6) to chromosome 17P", Human Molecular Genetics, 6:597-600 (1997)			
	H	Kelsell et al., "Localization of a Gene (CORD7) for a Dominant Cone-Rod Dystrophy to Chromosome 6q", Am. J. Hum. Genet., 63:274-279 (1998)			
	I	Stone et al., "Clinical Features of a Stargardt-Like Dominant Progressive Macular Dystrophy With Genetic Linkage to Chromosome 6q", Arch Ophthalmol., 112:765-772 (June 1994)			
	J	Nichols et al., "Butterfly-shaped pigment dystrophy of the fovea caused by a point mutation in codon 167 of the RDS gene", Nature, 3:202-207 (March 1993)			
	K	Weleber et al., "Phenotypic Variation Including Retinitis Pigmentosa, Pattern Dystrophy, and Fundus Flavimaculatus in a Single Family With a Deletion of Codon 153 or 154 of the Peripherin/RDS Gene, Arch Ophthalmol., 111:1531-1542 (November 1993)			
	L	Wells et al., "Mutations in the human retinal degeneration slow (RDS) gene can cause either retinitis pigmentosa or macular dystrophy", Nature Genet., 3:213-218 (March 1993)			
	M	Reig et al., "A point mutation in the RDS-peripherin gene in a Spanish family with central areolar choroidal dystrophy", Ophthalmic Genetics, 16:39-44 (1995)			
	N	Hyman et al., "Senile Macular Degeneration: A Case-Control Study", Am. J. Epidemiology, 118:213-227 (1983)			
↓	O	J. Donald M. Gass, "Drusen and Disciform Macular Detachment and Degeneration", Arch Ophthalmol., 90:206-217 (September 1973)			
AJ	P	Leppanen et al., "A Physical Map of the 6q14-q15 Region Harboring the Locus for the Lysosomal Membrane Sialic Acid Transport Defect", Genomics, 37:62-67 (1996)			

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			Group Art Unit	1646	
			Examiner Name		
Sheet	of	4	Attorney Docket Number	20430P	

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Examiner Initials*	Cite No.	Include name of the author, title, date, page(s), volume-issue number(s) and place of publication.
AJ	Q	Biervert et al., "A Potassium Channel Mutation in Neonatal Human Epilepsy", Science, 279:403-406 (January 1998)
	R	Singh et al., "A novel potassium channel gene, KCNQ2, is mutated in an inherited epilepsy of newborns", Nature Genetics, 18:25-29 (January 1998)
	S	Schroeder et al., "Moderate loss of function of cyclic-AMP-modulated KCNQ2/KCNQ3 K+ channels causes epilepsy", Nature, 396:687-690 (December 1998)
	T	Kubisch et al., "KCNQ4, a Novel Potassium Channel Expressed in Sensory Outer Hair Cells, Is Mutated in Dominant Deafness", Cell, 96: 437-446 (February 1999)
	U	Splawski et al., "Molecular Basis of the Long-QT Syndrome Associated with Deafness", New England J. Medicine, 336:1562-1567 (May 1997)
	V	Wang et al., "KCNQ2 and KCNQ3 Potassium Channel Subunits: Molecular Correlates of the M-Channel", Science, 282:1890-1893 (December 1998)
	W	Shi et al., "β Subunits Promote K+ Channel Surface Expression through Effects Early in Biosynthesis", Neuron, 16:843-852 (April 1996)
	X	Watson et al., "Molecular Biology of the Gene", Fourth Edition, The Benjamin/Cummings Publishing Co., Inc., page 226 (1987)
	Y	Cunningham and Wells, "High-Resolution Epitope Mapping of hGH-Receptor Interactions by Alanine-Scanning Mutagenesis", Science, 244:1081-1085 (June 1989)
	Z	Ioannou et al., "A new bacteriophage P1-derived vector for the propagation of large human DNA fragments", Nature Genetics, 6:84-89 (January 1994)
	AA	Gonzalez and Tsien, "Improved indicators of cell membrane potential that use fluorescence resonance energy transfer", Chemistry & Biology, 4:269-277 (1997)
	BB	Gonzalez and Tsien, "Voltage Sensing by Fluorescence Resonance Energy Transfer in Single Cells", Biophysical Journal, 69:1272-1280 (October 1995)
↓	CC	John Hodgson, "Receptor Screening and the Search for New Pharmaceuticals", Bio/Technology, 10:973-980 (September 1992)
AJ	DD	Hopp and Woods, "Prediction of protein antigenic determinants from amino acid sequences", Proc. Natl. Acad. Sci. USA, 78:3824-3828 (June 1981)

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AJ	EE	Jameson and Wolf, "The antigenic index: a novel algorithm for predicting antigenic determinants", CABIOS (Computer Applications in the Biosciences), 4:181-186 (1988)
	FF	Kohler and Milstein, "Continuous cultures of fused cells secreting antibody of predefined specificity", Nature, 256:495-497 (1975)
	GG	Thomas and Capecchi, "Site-Directed Mutagenesis by Gene Targeting in Mouse Embryo-Derived Stem Cells", Cell, 51:503-512 (November 1987)
	HH	Frohman and Martin, "Cut, Paste, and Save: New Approaches to Altering Specific Genes in Mice", Cell, 56:145-147 (January 1989)
	II	Mario R. Capecchi, "The New Mouse Genetics: Altering the Genome by Gene Targeting", Trends in Genetics, 5:70-76 (March 1989)
	JJ	Baribault and Kemler, "Embryonic Stem Cell Culture and Gene Targeting in Transgenic Mice", Mol. Biol. Med. 6:481-492 (1989)
	KK	Lin et al., "Recombination in mouse L cells between DNA introduced into cells and homologous chromosomal sequences", Proc. Natl. Acad. Sci. USA, 82:1391-1395 (March 1985)
	LL	Smithies et al., "Insertion of DNA sequences into the human chromosomal β-globin locus by homologous recombination", Nature, 317:230-234 (September 1985)
	MM	Thomas et al., "High Frequency Targeting of Genes to Specific Sites in the Mammalian Genome", Cell, 44:419-428 (February 1986)
	NN	Kim and Smithies, "Recombinant fragment assay for gene targeting based on the polymerase chain reaction", Nucleic Acids Research, IRL Press Ltd., Oxford, England, 16:8887-8903 (1988)
	OO	Kim et al., "Problems encountered in detecting a targeted gene by the polymerase chain reaction", Gene, 103:227-233 (1991)
	PP	Sedivy and Sharp, "Positive genetic selection for gene disruption in mammalian cells by homologous recombination", Proc. Natl. Acad. Sci. USA, 86:227-231 (January 1989)
↓ AJ	QQ	Mansour et al., "Disruption of the proto-oncogene <i>int-2</i> in mouse embryo-derived stem cells: a general strategy for targeting mutations to non-selectable genes", Nature, 336:348-352 (November 1988)
	RR	Mario R. Capecchi, "Altering the Genome by Homologous Recombination", Science, 244:1288-1292 (June 1989)

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